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# FY09 Scope of Work

WA Title: Impact of Green Building Products on Indoor Air Quality

# 1. Purpose

The overall objective of this project is to develop, demonstrate, and evaluate sustainable practices for indoor environments. Sustainable practices are decisions and actions that consider, minimize, and harmonize the impact of material and energy use on human health and the environment. Through integrated, multidisciplinary, and focused research, IEMB develops knowledge and tools that enable evaluation of sustainable practices for indoor environments. IEMB develops tools to characterize sources of indoor contaminants and investigates the relationships between sources of contaminants, the built environment and potential exposure to individual compounds and complex mixtures. For example, IEMB investigates the impact of green building products on indoor air quality and develops risk management options where green building practices or products may potentially improve or impair indoor quality.

# 2. Background

Rapidly increasing energy costs coupled with increasing market acceptance of "green" or sustainable residential building design has resulted in increased demand for sustainable building practices and "green" building products. However, sustainable "green" building practices (e.g., super insulated, tight buildings constructed with recycled or "natural" products) may inadvertently result in degraded indoor environmental quality or other downstream environmental challenges. As a component of "cradle to eradle" stewardship of materials and energy, there is a need to understand the impacts on the indoor environment of: emissions, sorption and re-emission of organic and inorganic compounds from "green" building materials (2) transport within the built environment, (3) efficacy of control technologies such as air and surface cleaning.

Key pollutants of concern include endocrine disrupting compounds such as brominated flame retardants, phthalates, and perfluorinated compounds associated with consumer products, neurotoxins such as elemental mereury released from the debris field of broken compact fluorescent light bulbs, and air toxics such as formaldehyde released and sorbed by numerous indoor materials and surfaces. Formaldehyde is one of the key toxic pollutants in the National Risk Management Research Laboratory (NRMRL) Indoor Air Strategic Plan. It is among the US Environmental Protection Agency (EPA) listed urban air hazardous air pollutants (HAPs) and one of the predominant VOCs emitted from building products. Primary emissions from materials and products as well as sorption and re-emission from surfaces are key factors that govern indoor concentrations. The sorption characteristics of wallboard significantly impact indoor concentrations of formaldehyde. However, this process is not well understood and the impact of "green" moisture resistant wallboard may alter the net sorption of formaldehyde, resulting in greater exposure to this known air toxic compound.

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There are three components of IEMB's research approach: (1) Develop source models that simulate emissions from green building products, (2) develop sorption/re-emission models for green building products, and (3) determine the reliability of source/sink models in full-scale indoor environments. The source emissions model parameters obtained from EPA's chamber tests will be applied to IAQ models to determine the impact of the use of "green" design building products on indoor concentrations organic and inorganic contaminants. Source and sink models and control strategies will be evaluated by studies conducted in APPCD's Research Test House (RTH), operated by the contractor. Specific tasks and the schedule for tasks to be conducted in the RTH will be described in amendments to this work assignment or described in other task-specific work assignments.

#### 3. Task Descriptions

The contractor shall conduct the following tasks:

The contractor shall: maintain the research house in ready mode for model evaluation or other studies as described in amendments to this work assignment or described in separate work assignments that utilize the research test house. Specifically, the contractor shall ensure that:

All miscellaneous and standard operating procedures (MOPs and SOPs) are accurate and up to date for contractor operated measurement or control systems. At a minimum, the contractor shall ensure that:

- The data acquisition system is functional
- · At least two temperature sensors and two RH sensors are functional
- The B&K Multi-gas Analyzer is calibrated for SF<sub>6</sub>
- The SF<sub>6</sub> dosing and sampling system is functional

Per section YY of the above referenced contract, the contractor shall maintain the instrumentation in the RTH to ensure that RTH can be utilized for specific research tasks within 30 days of notification through amendment to this or other work assignments.

The contractor shall provide technical input to QA test plans, addendums, technical reports, and manuscripts developed by EPA staff for and from specific experiments to be conducted in the research test house. Data gathering/manipulation shall not begin until the QAPP has been approved by the EPA QA Manager. The QA plan shall be developed according to the requirements in Attachment #1 to the Statement of Work. Specific experiments, schedules and deliverables will be described in amendments to this work assignment.

# 4. Reports

The contractor shall provide the EPA work assignment manager monthly progress reports as specified in the contract.

5. Schedule of Tasks, Reports, and Deliverables

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The contractor shall provide monthly reports of the RTH operational status. Reports and deliverables for other tasks, including new or revised MOPs or SOPs that are required to support QAPPs developed for specific research tasks to be conducted at the RTH, will be described in amendments to this work assignment.

# 6. Suggested Skills

This project will require contractor staff with the following skills: modification and adaptation of scientific apparatus to meet project objectives, sample collection and extraction, data processing and analysis, preparation, operation and maintenance of the RTH.

# 7. Special requirements

The contractor shall provide necessary health and safety procedures, documentation, and training to contractor staff to ensure safe conduct of the experiments at contractor controlled facilities.

# ATTACHMENT #1 TO THE STATEMENT OF WORK (SOW)

# NRMRL Quality Assurance (QA) Requirements

In accordance with EPA Order 5360.1 A2, conformance to ANSI/ASQC E4 must be demonstrated by submitting the quality documentation specified herein. All quality documentation shall be submitted to the Government for review. The Government will review and return the quality documentation, with comments, and indicate approval or disapproval. If the quality documentation is not approved, it must be revised to address all comments and shall be resubmitted to the Government for approval. Work involving environmental data collection, generation, use, or reporting shall not commence until the Government has approved the quality documentation. The quality documentation shall be submitted to the Government at least thirty (30) days prior to the beginning of any environmental data gathering or generation activity in order to allow sufficient time for review and revisions to be completed. After the Government has approved the quality documentation, the Contractor shall also implement it as written and approved by the Government. Any EPA-funded project/program may be subject to a QA audit.

#### TO BE SUBMITTED PRE-AWARD:

	NRMRL=s Quality System Specifications:			
	(1)	a description of the organization-s Quality System (QS) and information regarding how this QS is documented, communicated and implemented;		
).	(2)	an organizational chart showing the position of the QA function;		
	(3)	delineation of the authority and responsibilities of the QA function;		
	(4)	the background and experience of the QA personnel who will be assigned to the project; and		
	(5)	the organization—s general approach for accomplishing the QA specifications in the SOW.		
	Quali	ity Management Plan: prepared in accordance with R-2 - EPA Requirements for ty Management Plans (EPA/240/B-01/002) March, 2001, (www.cpa.gov/quality/qs-docs/r2-final.pdf		
то в	E SUB	MITTED POST-AWARD (mark all that apply):		
	NRM	RL=s Quality System Specifications:		
	(1)	a description of the organization=s Quality System (QS) and information regarding how this QS is documented, communicated and implemented;		

	(2)	an organizational chart showing the position of the QA function;		
	(3)	delineation of the authority and responsibilities of the QA function;		
	(4)	the background and experience of the QA personnel who will be assigned to the project; and		
	(5)	the organization=s general approach for accomplishing the QA specifications in the SOW.		
	Qualit	ty Management Plan: prepared in accordance with R-2 - EPA Requirements for y Management Plans (EPA/240/B-01/002) March, 2001, www.cpa.gov/quality/qs-docs/r2-final.pdf		
	with R	ory I or II Quality Assurance Project Plan (QAPP): prepared in accordance C-5 - EPA Requirements for QA Project Plans (EPA/240/B-01/003) March, 2001 www.epa.gov/quality/qs-docs/r5-final.pdf		
K	follow	egory III or IV QAPP: prepared in accordance with applicable sections of the owing NRMRL QAPP Requirements List(s) which is(are) included in this chment:		
	X	QAPP Requirements for Measurement Projects		
		QAPP Requirements for Secondary Data Projects		
		QAPP Requirements for Research Model Development and Application Projects		
		QAPP Requirements for Software Development Projects		
		QAPP Requirements for Method Development Projects		
		QAPP Requirements for Design, Construction, and Operation of Environmental Technology Projects		
ADDI	TIONA	AL QA RESOURCES:		
EPA=	s Quali	ity System Website: http://www.epa.gov/quality/		
EPA=	s Requ	irements and Guidance Documents: http://www.epa.gov/quality/qa_docs.htm		

(ATTACH APPROPRIATE QAPP REQUIREMENTS HERE)

# NRMRL QAPP REQUIREMENTS FOR MEASUREMENT PROJECTS

**GENERAL REQUIREMENTS:** Include cover page, distribution list, approvals, and page numbers.

#### 0. COVER PAGE

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Include the Division/Branch, project title, revision number, EPA technical lead, QA category, organization responsible for QAPP preparation, and date.

#### 1. PROJECT DESCRIPTION AND OBJECTIVES

- 1.1 Describe the process and/or environmental system to be evaluated.
- 1.2 State the purpose of the project and list specific project objective(s).

#### 2. ORGANIZATION AND RESPONSIBILITIES

- 2.1 Identify all project personnel, including QA, and related responsibilities for each participating organization, as well as their relationship to other project participants.
- 2.2 Include a project schedule that includes key milestones.

#### 3. SCIENTIFIC APPROACH

- 3.1 Describe the sampling and/or experimental design that will be used to generate the data needed to evaluate the projective objective(s). A description of the design should include the types and numbers of samples (including QC and reserve samples), the design of the sampling network, sample locations and frequencies, and the rationale for the design.
- 3.2 Identify the process measurements (e.g., flow rate, temperature) and specific target analyte(s) for each sample type.
- 3.3 Describe the general approach and the test conditions for each experimental phase.

#### 4. SAMPLING PROCEDURES

- 4.1 Describe any known site-specific factors that may affect sampling procedures as well as all site preparation (e.g., sampling device installation, sampling port modifications, achievement of steady-state) needed prior to sampling.
- 4.2 Describe or reference each sampling procedure (including a list of equipment needed and the calibration of this equipment as appropriate) to be used. Include procedures for homogenizing, compositing, or splitting of samples, as applicable.
- 4.3 Provide a list of sample containers, sample quantities to be collected, and the sample amount required for each analysis, including QC sample analysis.
- 4.4 Specify sample preservation requirements (e.g., refrigeration, acidification, etc.) and holding times.
- 4.5 Describe the method for uniquely numbering each sample.

4.6 Describe procedures for packing and shipping samples, including procedures to avoid cross-contamination, and provisions for maintaining chain-of-custody (e.g., custody seals and records), as applicable.

#### 5. MEASUREMENT PROCEDURES

- 5.1. Describe in detail or reference each process measurement or analytical method to be used. If applicable, identify modifications to EPA-approved or similarly validated methods.
- 5.2. If not provided in Section 5.1 or the referenced method, include specific calibration procedures, including linearity checks and initial and continuing calibration checks.

# 6. QUALITY METRICS (QA/QC CHECKS)

- 6.1. For each process measurement and analytical method, identify the required QC checks (e.g., blanks, control samples, duplicates, matrix spikes, surrogates), the frequencies for performing these checks, associated acceptance criteria, and corrective actions to be performed if acceptance criteria are not met.
- 6.2. Any additional project-specific QA objectives (e.g., completeness, mass balance) shall be presented, including acceptance criteria.

# 7. DATA ANALYSIS, INTERPRETATION, AND MANAGEMENT

- 7.1 Identify the data reporting requirements, including data reduction procedures specific to the project and applicable calculations and equations.
- 7.2 Describe data validation procedures used to ensure the reporting of accurate project data.
- 7.3 Describe how the data will be summarized or analyzed (e.g., qualitative analysis, descriptive or inferential statistics) to meet the project objective(s).
  - 7.3.1 If descriptive statistics are proposed, state what tables, plots, and/or statistics (e.g., mean, median, standard error, minimum and maximum values) will be used to summarize the data.
  - 7.3.2 If an inferential method is proposed, indicate whether the method will be a hypothesis test, confidence interval, or confidence limit and describe how the method will be performed.
- 7.4 Describe data storage requirements for both hard copy and electronic data.

#### 8. REPORTING

- 8.1 List and describe the deliverables expected from each project participant responsible for field and/or analytical activities.
- 8.2 Specify the expected final product(s) that will be prepared for the project (e.g., journal article, final report).

#### 9. REFERENCES

Provide references either in the body of the text as footnotes or in a separate section.